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Introduction

DNA barcoding aims at using a standard 647bp fragment of the mitochondrial cytochrome c oxidase subunit I (COI) gene to identify species. Unfortunately, DNA of museum specimens is often degraded. Therefore, smaller fragments of the COI gene are used as mini-barcodes. Here, we compared the identification success of the standard (BC) and mini-barcode (mini-BC) fragment in a group of cosmopolitan raptors.

Material & methods

- 25 European and African *Accipiter* species
- 140 specimens

- three datasets: A: 25 species, 291bp, B: 19 species, 647bp, C: 19 species, 291bp



Century old photo of *Accipiter* (then *Astur*) *castanilius*

- intra- and interspecific sequence divergence: K2P distances
- Identification based on:
 - best match (BM) criterion,
 - best close match (BCM) criterion and
 - character based identification
- thresholds using intra- and interspecific comparisons:
 - 10X mean intraspecific distance
 - best compromise (BCo)



A. toussenellii females, Kinshasa MRAC 11995 & 7731 A.2

Examples of *Accipiter* skins used in this study



A. canescens females, Ualie MRAC 54432 & 47473

Results

- thresholds: 10X: 3-9% - 5.3% - BCo: 2.8% - 3.0%
- identification success: standard BC: 90% - mini-BC: 84%
- BM and BCM criteria perform equally well

a BM

dataset	BCTh threshold	correct	ambiguous	incorrect	no match closer than threshold
A	3.00%	124 (84.35%)	19 (12.92%)	4 (2.72%)	-
B	2.80%	90 (90.0%)	6 (6.0%)	4 (4.0%)	-
C	3.00%	90 (90.0%)	5 (5.0%)	5 (5.0%)	-

b BCM

dataset	BCTh threshold	correct	ambiguous	incorrect	no match closer than threshold
A	3.00%	124 (84.35%)	18 (12.24%)	3 (2.04%)	2 (1.36%)
B	2.80%	90 (90.0%)	5 (5.0%)	4 (4.0%)	1 (1.0%)
C	3.00%	90 (90.0%)	5 (5.0%)	4 (4.0%)	1 (1.0%)

c BM

dataset	10 X threshold	correct	ambiguous	incorrect	no match closer than threshold
A	4.80%	124 (84.35%)	19 (12.92%)	4 (2.72%)	-
B	5.30%	90 (90.0%)	5 (5.0%)	5 (5.0%)	-
C	3.90%	90 (90.0%)	6 (6.0%)	4 (4.0%)	-

d BCM

dataset	10 X threshold	correct	ambiguous	incorrect	no match closer than threshold
A	4.80%	124 (84.35%)	18 (12.24%)	3 (2.04%)	2 (1.36%)
B	5.30%	90 (90.0%)	5 (5.0%)	4 (4.0%)	1 (1.0%)
C	3.90%	90 (90.0%)	5 (5.0%)	4 (4.0%)	1 (1.0%)

Table: identification successes and threshold values for the different methods and criteria used.

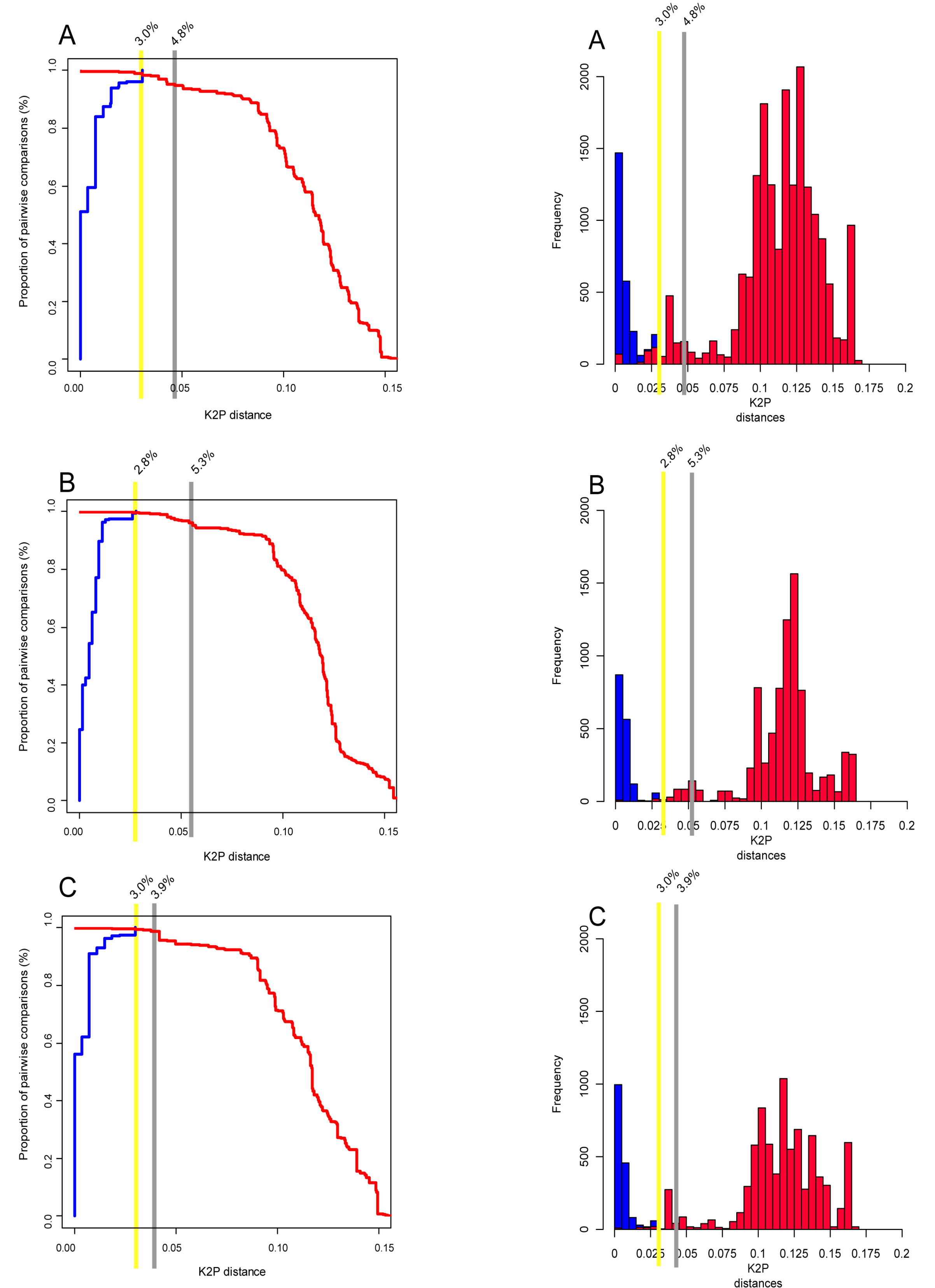


Figure 1: Cumulative frequency graphs of intra (blue) and interspecific distances (red) for the three datasets A, B & C

Figure 2: K2P distance distribution histograms with intra (blue) and interspecific distances (red) for the three datasets A, B & C

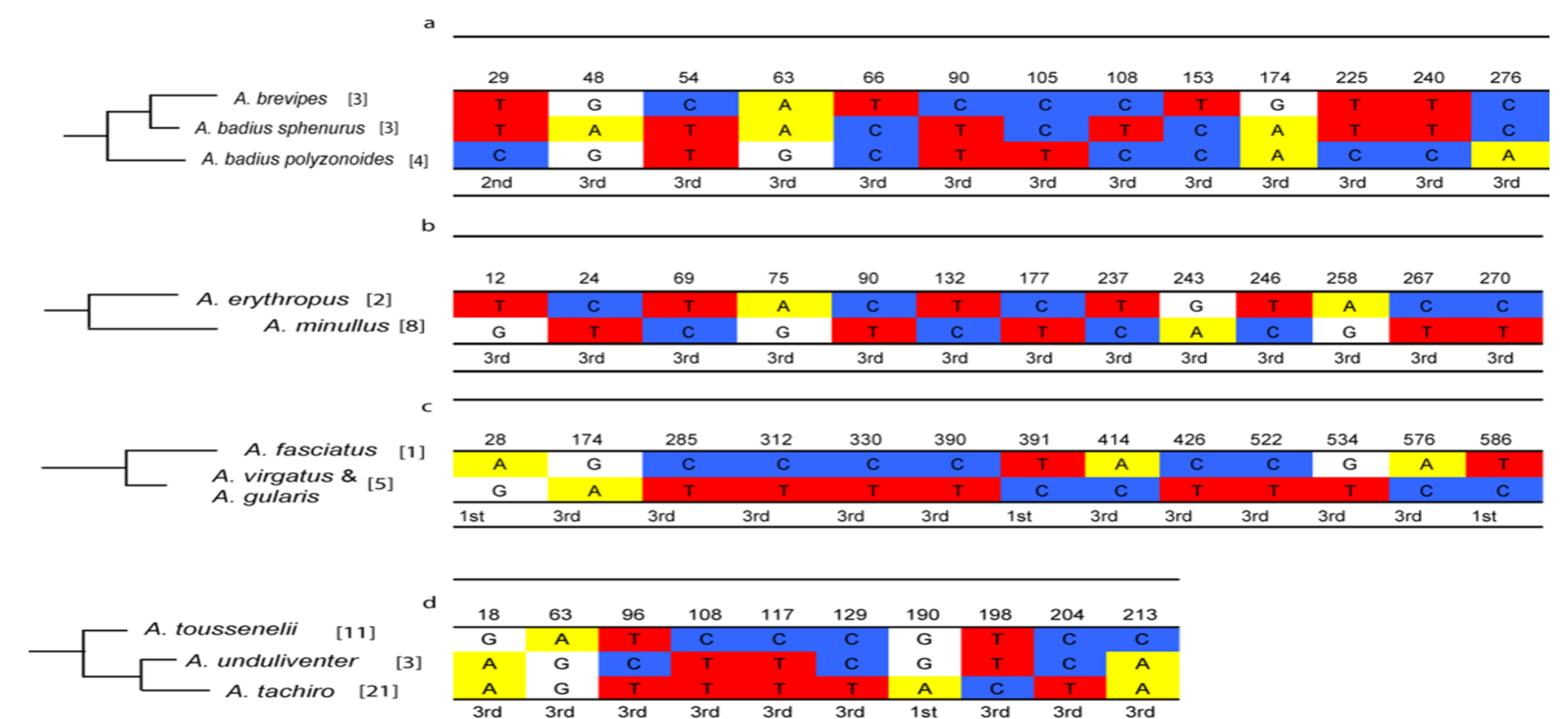


Figure 3: Character state analysis for four closely related groups of *Accipiter* species

Conclusions

- 19 of the *Accipiter* species studied can be identified unambiguously using DNA barcoding
- six species (three species pairs) can not be identified using DNA barcoding
- the mini-BC performs slightly worse than the standard BC